

Amendments to the Claims

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1-19. (Previously cancelled)

20. (Previously Added) A method of increasing cerebral bioavailability of a physiologically active composition in an individual comprising administering an NO-increasing agent or agents to the individual in need of enhanced drug delivery, wherein said agent is administered substantially contemporaneously with the physiologically active composition whose delivery is to be enhanced and said agent or agents increases the production of NO by preexisting ecNOS.

21. (Previously Added) A method of increasing cerebral bioavailability of a physiologically active composition in an individual comprising introducing the composition into the blood stream of the individual substantially contemporaneously with a blood flow enhancing amount of L-arginine.

22. (Previously Added) A method of increasing cerebral bioavailability of a physiologically active composition in an individual according to claim 20 comprising introducing the composition into the blood stream of the individual substantially contemporaneously with a blood flow enhancing amount of an agent which increases the production of NO by preexisting ecNOS and at least one other NO-increasing agent.

23. (Amended) The method according to claim [s20 and] 22, [further] wherein the agent which increases the production of NO by preexisting ecNOS is selected from the group consisting of L-arginine, NADPH, and tetrahydrobiopterin.

24. (Previously Added) The method according to claim 23, further wherein the agent which increases the production of NO by preexisting ecNOS is L-arginine.

25. (Previously Added) The method according to claim 22, wherein the agent which increases the production of NO by preexisting ecNOS is L-arginine and the at least one other NO-increasing agent is a different an agent which increases the production of NO by preexisting ecNOS.

26. (Previously Added) The method according to claim 22, wherein the agent which increases the production of NO by preexisting ecNOS is L-arginine and the at least one other NO-increasing agent is a non-ecNOS NO-generating system.

27. (Amended) The method according to any one of claims 20 to 22 wherein the individual in need of enhanced drug delivery has experienced, is experiencing, or has an [is-at] abnormally elevated risk of experiencing an ischemic stroke.

28. (Previously Added) The method according to any one of claims 20 to 22 wherein the physiologically active composition has a site of action in the brain.

29. (Amended) A pharmaceutical composition [for increasing cerebral bioavailability of a physiologically active composition in an individual] comprising a blood-flow enhancing amount of an NO-increasing agent or agents which increases the production of NO by preexisting ecNOS and [the] a physiologically active composition.

30. (Previously Added) The composition according to claim 29, wherein the NO-increasing agent comprises an agent which increases the production of NO by preexisting ecNOS and at least one other NO-increasing agent.

31. (Previously Added) The composition according to claim 29 or claim 30, wherein the agent which increases the production of NO by preexisting ecNOS is selected from the group consisting of L-arginine, NADPH, and tetrahydrobiopterin.

32. (Previously Added) The composition according to claim 29 or claim 30, wherein the agent which increases the production of NO by preexisting ecNOS is L-arginine.

33. (Amended) The composition according to claim 30, comprising L-arginine and at least one other NO-increasing agent which is a different [an] agent which increases the production of NO by preexisting ecNOS.

34. (Previously Added) The composition according to claim 30, comprising L-arginine and at least one NO-increasing agent which is a non-ecNOS NO-generating system.

35. (Amended) [The composition according to claim 29 or claim 30, wherein the physiologically active composition has]

A pharmaceutical composition comprising a blood-flow enhancing amount of an NO-increasing agent or agents which increases the production of NO by preexisting ecNOS and a physiologically active composition having a site of action in the brain.